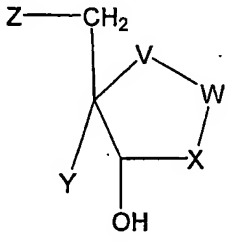
Amendments to the Claims:

Please cancel Claim 22 without prejudice or disclaimer, and amend Claims 1, 25 and 27 as set forth below.

1. (Currently amended) A compound of the formula (I):



(I)

wherein:

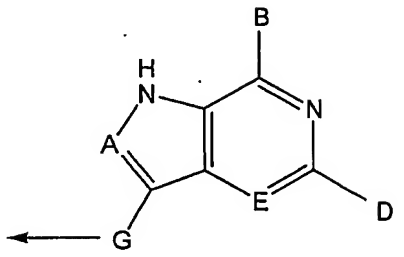
V is selected from CH_2 and NH, and W is NR^1; or V is NR^1, and W is selected from CH_2 and NH;

X is selected from CH_2 and CHOH in the R or S-configuration;

Y is selected from hydrogen, halogen and hydroxy, except where V is selected from NH and NR^1 then Y is hydrogen;

Z is selected from hydrogen, halogen, hydroxy, SQ, OQ and Q, where Q is an ~~optionally substituted~~ alkyl, aralkyl or aryl group, where the aryl group is optionally substituted;

R^1 is a radical of the formula (II)



(II)

A is selected from CH and CR, where R is selected from halogen, ~~optionally substituted~~ alkyl, aralkyl or aryl, OH, NH_2, NHR^3, NR^3R^4 and SR^5, where R^3, R^4 and R^5 are each ~~optionally substituted~~ alkyl, aralkyl or aryl groups;

B is selected from OH, NH_2, NHR^6, SH, hydrogen and halogen, where R^6 is an ~~optionally substituted~~ alkyl, aralkyl or aryl group;

D is selected from OH, NH_2, NHR^7, hydrogen, halogen and SCH_3, where R^7 is an ~~optionally substituted~~ alkyl, aralkyl or aryl group;

E is ~~selected from~~ N ~~and CH~~;

G is selected from CH_2 and NH, or G is absent, provided that where W is NR^1 and G is NH then V is CH_2, and provided that where V is NR^1 and G is NH then W is CH_2,

or a tautomer thereof, or a pharmaceutically acceptable salt thereof.

2. (Previously presented) A compound as claimed in claim 1, where Z is selected from hydrogen, halogen, hydroxy, SQ and OQ.

3. (Previously presented) A compound as claimed in claim 1, where V is CH_2.

4. (Previously presented) A compound as claimed in claim 1, where X is CH_2.

5. (Previously presented) A compound as claimed in claim 1, where G is CH_2.

6. (Previously presented) A compound as claimed in claim 1, where Z is OH.

7. (Previously presented) A compound as claimed in claim 1, where Z is SQ.

8. (Previously presented) A compound as claimed in claim 1, where Z is Q.

9. (Previously presented) A compound as claimed in claim 1, where W is NR^1.

10. (Canceled)

11. (Previously presented) A compound as claimed in claim 1, where W is selected from NH and NR^1 and X is CH_2.

12. (Previously presented) A compound as claimed in claim 1, where V, X and G are all CH_2, Z is OH and W is NR^1.

13. (Previously presented) A compound as claimed in claim 1, where V, X and G are all CH_2, Z is SQ and W is NR^1.

14. (Previously presented) A compound as claimed in claim 1, where Y is hydrogen.

15. (Previously presented) A compound as claimed in claim 1, where Y is hydroxy.

16. (Previously presented) A compound as claimed in claim 1, where B is hydroxy.

17. (Previously presented) A compound as claimed in claim 1, where B is NH_2.

18. (Previously presented) A compound as claimed in claim 1, where A is CH.

19. (Canceled)

20. (Previously presented) A compound as claimed in claim 1, where D is H.

21. (Previously presented) A compound as claimed in claim 1, where D is NH_2.

22. (Canceled)

23. (Previously presented) A compound as claimed in claim 1, which is: (3R,4R)-1-[(9-deazahypoxanthin-9-yl)methyl]-3-hydroxy-4-(hydroxymethyl)pyrrolidine; (3R,4R)-1-[(9-Deazaadenin-9-yl)methyl]-3-hydroxy-4-(hydroxylmethyl)pyrrolidine; (3R,4S)-1-[(9-deazaadenin-9-yl)methyl]-3-hydroxy-4-(2-phenylethyl)pyrrolidine;

(3S,4R)-1-[(9-deazahypoxanthin-9-yl)methyl]-3,4-dihydroxy-4-methylthiomethyl pyrrolidine;

(3R,4S)-1-[(9-deazahypoxanthin-9-yl)methyl]-3-hydroxy-4-(methylthiomethyl) pyrrolidine;

N-(9-Deazahypoxanthin-9-yl)-1,4-dideoxy-1,4-imino-D-ribitol;

N-(9-deazahypoxanthin-9-yl)methyl-1,4-dideoxy-1,4-imino-D-ribitol;

(3R,4S)-1-[(9-deazaadenin-9-yl)methyl]-3-hydroxy-4-(methylthiomethyl)pyrrolidine;

(3R,4S)-1[(9-deazaadenin-9-yl)methyl]-3-hydroxy-4-(benzylthiomethyl)pyrrolidine;

(3R,4R)-1-[(9-deazaguanin-9-yl)methyl]-3-hydroxy-4-(hydroxymethyl)pyrrolidine;

(3R,4S)-1-[(9-deazaadenin-9-yl)methyl]-3-hydroxy-4-(4-chlorophenylthiomethyl) pyrrolidine;

(3R,4R)-1-[(6-chloro-9-deazapurin-9-yl)methyl]-3-hydroxy-4-(hydroxymethyl) pyrrolidine;

(3R,4R)-1-[(6-azido-9-deazapurin-9-yl)methyl]-3-hydroxy-4-(hydroxymethyl) pyrrolidine; or

(3R,4R)-1-[(9-deezaadenin-9-yl)methyl]-3-acetoxy-4-(acetoxymethyl)pyrrolidine;

or a pharmaceutically acceptable salt thereof.

24. (Previously presented) A pharmaceutical composition comprising a pharmaceutically effective amount of a compound as claimed in claim 1 and a carrier.

25. (Currently amended) A method of treating a subject ~~having a disease or condition in which it is desirable to inhibit purine phosphoribosyltransferase, purine nucleoside phosphorylase, 5'-methylthio adenosine phosphorylase, 5'-methylthioadenosine nucleosidase and/or nucleoside hydrolase~~ comprising administering a compound as claimed in claim 1 to the subject ~~in an amount effective to~~

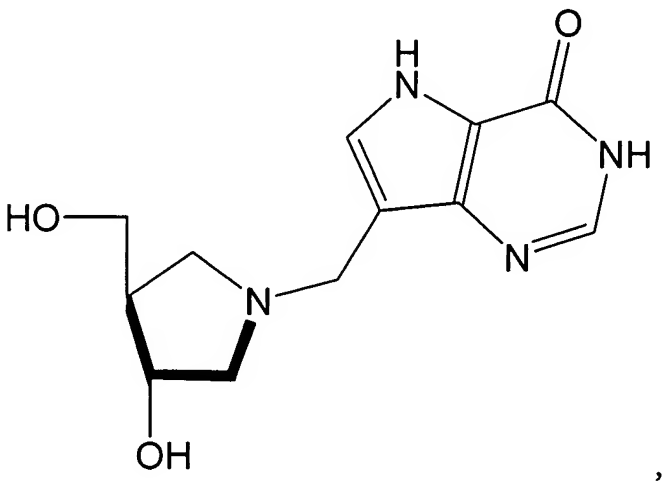
~~inhibit purine phosphoribosyltransferase, purine nucleoside phosphorylase, 5'-methylthio adenosine phosphorylase, 5'-methylthioadenosine nucleosidase and/or nucleoside hydrolase~~, wherein the subject has ~~disease or condition is~~ a cancer, a bacterial infection, a protozoal infection, a T-cell mediated disease or a transplant rejection.

26. (Canceled)

27. (Currently amended) The method of claim 25, where the T-cell mediated disease is psoriasis ~~or arthritis~~.

28. (Canceled)

29. (Previously presented) The compound of claim 1 having the structure:



,

or a pharmaceutically acceptable salt thereof.

30. (Previously presented) The compound of claim 1, wherein the salt is derived from an acid selected from the group consisting of hydrochloric acid, sulphuric acid, phosphoric acid, acetic acid, lactic acid, fumaric acid, succinic acid, tartaric acid, gluconic acid, citric acid, methanesulfonic acid and p-toluenesulfonic acid.

31. (Previously presented) The compound of claim 29, wherein the salt is a hydrochloride salt.